SALT AND STONE SPF 30 SUNSCREEN STICK- zinc oxide stick Salt and Stone

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Salt & Stone SPF 30 Sunscreen stick

DRUG FACTS

Active Ingredients

Zinc Oxide (Non-Nano) 22.5%

Purpose

Sunscreen

Uses

- Hepls prevent sunburn.
- If used as directed with other sun protection measures (see directions), decreases the risk of skin cancer and early skin aging caused by the sun.

Warnings

For external use only.

Do not use

on damaged or broken skin.

When using this product,

keep out of eyes. Rinse with water to remove.

If rash occurs

stop use and see a doctor.

Keep out of reach of children.

If product is swallowed, get medical help or contact a poison control center right away.

Directions

- Apply liberally 15 minutes before sun exposure.
- Reapply:
- After 80 minutes of swimming or sweating,

- Immediately after towel drying.
- At least every two (2) hours.
- **Sun Protection Measures** Spending time in the sun increases the risk of skin cancer and early skin aging. To decrease the risk, regularly use: a suscreen with a broad-spectrum measures including:
- Limiting time in the sun, especially from 10AM 2PM.
- Wear long-sleeved shirts, pants, hats, and sunglasses.
- Children under 6 months: Ask a doctor.

Other Information

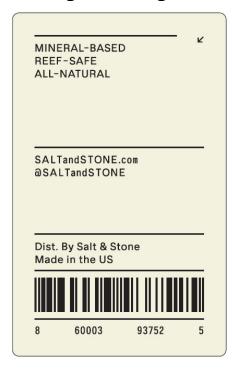
Protect this product from excess heat and direct sun.

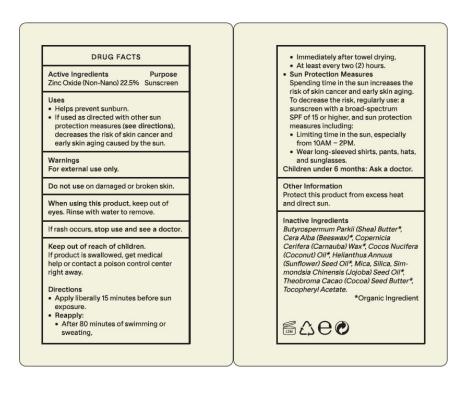
Inactive Ingredients

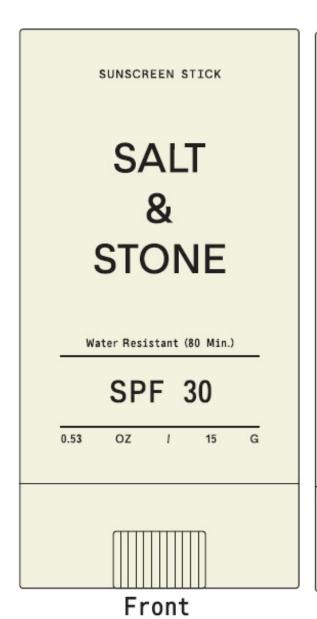
Butyrospermum Parkii (Shea) Butter, Cera Alba (Beeswax)*, Copernicia Cerifera (Carnauba) Wax,* Cocos Nucifera (Coconut) Oil,* Helianthus Annuus (Sunflower) Seed Oil,* Mica, Silica, Simmondsia Chinensis (Jojoba) Seed Oil,* Theobroma Cacao (Cocoa) Seed Butter, Tocopheryl Acetate.

*Organic Ingredient

Package Labeling:









SALT AND STONE SPF 30 SUNSCREEN STICK

zinc oxide stick

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71585-141
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC CATION - UNII:13S1S8SF37)	ZINC CATION	225 mg in 1 g

Inactive Ingredients	
Ingredient Name	Strength
SHEA BUTTER (UNII: K49155WL9Y)	

YELLOW WAX (UNII: 2ZA36H0S2V)	
CARNAUBA WAX (UNII: R12CBM0EIZ)	
COCONUT OIL (UNII: Q9L0O73W7L)	
SUNFLOWER OIL (UNII: 3W1JG795YI)	
MICA (UNII: V8A1AW0880)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
JOJOBA OIL (UNII: 724GKU717M)	
COCOA BUTTER (UNII: 5120YT1CRR)	
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

F	Packaging			
#	tem Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71585-141- 00	15 g in 1 TUBE; Type 0: Not a Combination Product	09/22/2021	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part352	09/22/2021	

Labeler - Salt and Stone (080683697)

Revised: 11/2021 Salt and Stone